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IN THE UNITED STATES DISTRICT COURT  
FOR THE NORTHERN DISTRICT OF CALIFORNIA  
SAN JOSE DIVISION

AMY MAXWELL, individually and on  
behalf of all others similarly situated,

Plaintiff,

v.

UNILEVER UNITED STATES, INC.,  
UNILEVER SUPPLY CHAIN, INC.,  
and PEPSICO, INC.,

Defendants.

Case No.

**CLASS ACTION AND REPRESENTATIVE  
ACTION**

**COMPLAINT FOR DAMAGES,  
EQUITABLE AND INJUNCTIVE RELIEF**

**JURY TRIAL DEMANDED**

Plaintiff, through her undersigned attorneys, brings this lawsuit against Defendants as to her own acts upon personal knowledge, and as to all other matters upon information and belief. In order to remedy the harm arising from Defendants' illegal conduct, which has resulted in unjust profits, Plaintiff brings this action on behalf of a class of California consumers who purchased Lipton Tea products within the last four years.

**INTRODUCTION**

1. Every day, millions of Americans purchase and consume packaged foods. Identical federal and California laws require truthful, accurate information on the labels of

1 packaged foods. This case is about a company that flouts those laws, before and after receiving a  
 2 warning letter from the FDA. The law, however, is clear: misbranded food cannot legally be  
 3 manufactured, held, advertised, distributed or sold. Misbranded food is worthless as a matter of  
 4 law, and purchasers of misbranded food are entitled to a refund of their purchase price.

5 2. Unilever is a multinational corporation with 400 brands, including Lipton Tea.  
 6 Unilever's website claims that "[o]n any given day, two billion people use our products." Lipton  
 7 employs "more than 80,000 people." According to Unilever, "tea is the second most widely-  
 8 consumed beverage on earth, behind water." In the U.S., Unilever markets Lipton Tea under  
 9 more than twelve labels, including Lipton Regular Tea, Lipton Cold Brew Iced Tea Bags, Lipton  
 10 Green Tea Purple Acai and Blueberry, Lipton Regular Family Size Iced Tea Bags, Lipton Lemon  
 11 Iced Tea Mix, Lipton Diet Lemon Ice Tea Mix, Lipton Green Tea, Lipton Mandarin Mango  
 12 Green Tea To Go, Lipton Herbal Chamomile Tea, Lipton Harvest Strawberry & Passion Fruit,  
 13 Lipton Orange Spice Tea and Lipton Ready to Drink PureLeaf Tea.

14 <http://www.unileverusa.com/brands/foodbrands/lipton/index.aspx>

15 3. Unilever recognizes that health claims drive sales, and actively promotes the health  
 16 benefits of Lipton Tea, including flavonoids:

17 Made from real tea leaves, many Lipton teas contain tea flavonoids. The flavonoid  
 18 content per serving can be found on all Lipton tea packages with the Tea Goodness  
 19 seal which signals that the tea contains a specific level of tea flavonoids.  
 20 Flavonoids are dietary compounds found in tea, wine, cocoa, fruit and vegetables.  
 21 They contribute significantly to taste and color, and possibly help maintain certain  
 22 normal, healthy body functions. A diet rich in flavonoids is generally associated  
 23 with helping maintain normal healthy heart function.

24 <http://www.unileverusa.com/brands/foodbrands/lipton/index.aspx>

25 4. On the Lipton Tea website, Unilever goes even further in promoting the health  
 26 benefits of Lipton Tea, specifically focusing on tea flavonoids:

27 Studies suggest that drinking black or green tea may help maintain normal, healthy  
 28 heart function as part of a diet that is consistent with dietary guidelines. Research  
 suggests that drinking 2 to 3 cups per day of black or green tea may help support  
 normal, healthy vascular function. The mechanism behind this effect has yet to be  
 fully demonstrated, but research suggests that tea flavonoids may be responsible.

[http://www.lipton.com/tea\\_health/healthy\\_diet/index.aspx](http://www.lipton.com/tea_health/healthy_diet/index.aspx)

1           5.       Unilever makes health nutrient claims directly on packages of Lipton Tea. For  
2 example, the package front panel of certain Lipton Tea products bears the “AOX Naturally  
3 Protective Antioxidants” label. The back panel further touts the “protective flavonoid  
4 antioxidants” and “flavonoid content” of Lipton Tea, by comparing Lipton Tea to “selected  
5 beverages and fruits,” including orange juice, broccoli, cranberry juice and coffee. Lipton  
6 Decaffeinated Tea Bags, 72 count.

7           6.       In promoting the health benefits of its products, including Lipton Tea, Unilever  
8 adopted “Global Principles for Responsible Food and Beverage Marketing.” These Global  
9 Principles apply to “all of Unilever’s food and beverage marketing activities and  
10 communications,” and include the following provisions:

11               These marketing activities and communications include but are not limited to packaging  
12               and labeling . . .

13               Marketing communications must comply with all relevant laws/regulations in the local  
14               country . . .

15               All food and beverage marketing communications must be truthful and not misleading.

16               [www.unileverusa.com/Images/30370 Global Principles A5 PDF-2 tcm23-48998.pdf](http://www.unileverusa.com/Images/30370_Global_Principles_A5_PDF-2_tcm23-48998.pdf)

17           7.       If a manufacturer is going to make a claim on a food label, the label must meet  
18 certain legal requirements that help consumers make informed choices and ensure that they are  
19 not misled. As described more fully below, Defendants have made, and continue to make, false  
20 and deceptive claims in violation of federal and California laws that govern the types of  
21 representations that can be made on food labels. These laws recognize that reasonable consumers  
22 are likely to choose products claiming to have a health or nutritional benefit over otherwise  
23 similar food products that do not claim such benefits.

24           8.       Identical federal and California laws regulate the content of labels on packaged  
25 food. The requirements of the federal Food Drug & Cosmetic Act (“FDCA”) were adopted by  
26 the California legislature in the Sherman Food Drug & Cosmetic Law (the “Sherman Law”).  
27 California Health & Safety Code § 109875, et seq. Under FDCA section 403(a), food is  
28 “misbranded” if “its labeling is false or misleading in any particular,” or if it does not contain  
certain information on its label or in its labeling. 21 U.S.C. § 343(a).

9. Under the FDCA, the term “false” has its usual meaning of “untruthful,” while the term “misleading” is a term of art. Misbranding reaches not only false claims, but also those claims that might be technically true, but still misleading. If any one representation in the labeling is misleading, then the entire food is misbranded, nor can any other statement in the labeling cure a misleading statement. “Misleading” is judged in reference to “the ignorant, the unthinking and the credulous who, when making a purchase, do not stop to analyze.” *United States v. El-O-Pathic Pharmacy*, 192 F.2d 62, 75 (9<sup>th</sup> Cir. 1951). Under the FDCA, it is not necessary to prove that anyone was actually misled.

10. On August 23, 2010, the FDA sent a warning letter to Unilever, informing Unilever of its failure to comply with the requirements of the FDCA and its regulations (the “FDA Warning Letter,” attached hereto as Exhibit 1). The FDA Warning Letter stated, in pertinent part:

#### **Unauthorized Nutrient Content Claims**

Under section 403(r)(1)(A) of the Act [21 U.S.C. 343(r)(1)(A)], a claim that characterizes the level of a nutrient which is of the type required to be in the labeling of the food must be made in accordance with a regulation promulgated by the Secretary (and, by delegation, FDA) authorizing the use of such a claim. The use of a term, not defined by regulation, in food labeling to characterize the level of a nutrient misbrands a product under section 403(r)(1)(A) of the Act.

Nutrient content claims using the term “antioxidant” must also comply with the requirements listed in 21 CFR 101.54(g). These requirements state, in part, that for a product to bear such a claim, an RDI must have been established for each of the nutrients that are the subject of the claim (21 CFR 101.54(g)(1)), and these nutrients must have recognized antioxidant activity (21 CFR 101.54(g)(2)). The level of each nutrient that is the subject of the claim must also be sufficient to qualify for the claim under 21 CFR 101.54(b), (c), or (e) (21 CFR 101.54(g)(3)). For example, to bear the claim “high in antioxidant vitamin C,” the product must contain 20 percent or more of the RDI for vitamin C under 21 CFR 101.54(b). Such a claim must also include the names of the nutrients that are the subject of the claim as part of the claim or, alternatively, the term “antioxidant” or “antioxidants” may be linked by a symbol (e.g., an asterisk) that refers to the same symbol that appears elsewhere on the same panel of the product label, followed by the name or names of the nutrients with recognized antioxidant activity (21 CFR 101.54(g)(4)). The use of a nutrient content claim that uses the term “antioxidant” but does not comply with the requirements of 21 CFR 101.54(g) misbrands a product under section 403(r)(2)(A)(i) of the Act.

Your webpage entitled “Tea and Health” and subtitled “Tea Antioxidants” includes the statement, “LIPTON Tea is made from tea leaves rich in naturally protective

1 antioxidants." The term "rich in" is defined in 21 CFR 101.54(b) and may be used  
2 to characterize the level of antioxidant nutrients (21 CFR 101.54(g)(3)). However,  
3 this claim does not comply with 21 CFR 101.54(g)(4) because it does not include  
4 the nutrients that are the subject of the claim or use a symbol to link the term  
"antioxidant" to those nutrients. Thus, this claim misbrands your product under  
section 403(r)(2)(A)(i) of the Act.

5 This webpage also states that "tea is a naturally rich source of antioxidants." The  
6 term "rich source" characterizes the level of antioxidant nutrients in the product  
7 and, therefore, this claim is a nutrient content claim (see section 403(r)(1) of the  
8 Act and 21 CFR 101.13(b)). Even if we determined that the term "rich source"  
9 could be considered a synonym for a term defined by regulation (e.g., "high" or  
10 "good source"), nutrient content claims that use the term "antioxidant" must meet  
11 the requirements of 21 CFR 101.54(g). The claim "tea is a naturally rich source of  
antioxidants" does not include the nutrients that are the subject of the claim or use  
a symbol to link the term "antioxidant" to those nutrients, as required by 21 CFR  
101.54(g)(4). Thus, this claim misbrands your product under section  
403(r)(2)(A)(i) of the Act.

12 The product label back panel includes the statement "packed with protective  
13 FLAVONOID ANTIOXIDANTS." The term "packed with" characterizes the level  
14 of flavonoid antioxidants in the product; therefore, this claim is a nutrient content  
15 claim (see section 403(r)(1) of the Act and 21 CFR 101.13(b)). Even if we  
16 determined that the term "packed with" could be considered a synonym for a term  
17 defined by regulation, nutrient content claims that use the term "antioxidant" must  
18 meet the requirements of 21 CFR 101.54(g). The claim "packed with  
FLAVONOID ANTIOXIDANTS" does not comply with 21 CFR 101.54(g)(1)  
because no RDI has been established for flavonoids. Thus, this unauthorized  
nutrient content claim causes your product to be misbranded under section  
403(r)(2)(A)(i) of the Act.

19 The above violations are not meant to be an all-inclusive list of deficiencies in  
20 your products or their labeling. It is your responsibility to ensure that all of your  
21 products are in compliance with the laws and regulations enforced by FDA. You  
22 should take prompt action to correct the violations. Failure to promptly correct  
these violations may result in regulatory actions without further notice, such as  
seizure and/or injunction.

23 We note that your label contains a chart entitled "Flavonoid Content of selected  
24 beverages and foods." The chart appears to compare the amounts of antioxidants in  
25 your product with the amount of antioxidants in orange juice, broccoli, cranberry  
26 juice and coffee. However, the information provided may be misinterpreted by the  
27 consumer because although the chart is labeled, in part, "Flavonoid Content," the  
y-axis is labeled "AOX"; therefore, the consumer might believe that the chart is  
stating the total amount of antioxidants rather than specifically measuring the  
amount of flavonoids in the product.

28 <http://www.fda.gov/ICECI/EnforcementActions/WarningLetters/ucm224509.htm>

11. In response to the FDA Warning letter, Unilever modified its Lipton web site and its packaging by removing some of the most outlandish claims of health and therapeutic benefits that FDA had found in violation of law. However, there are several unlawful statements on Lipton's web site that remain, including but not limited to: "Flavonoids are dietary compounds found in tea, wine, cocoa, fruit and vegetables. They contribute significantly to taste and color, and possibly help maintain certain normal, healthy body functions. A diet rich in flavonoids is generally associated with helping maintain normal, healthy heart function." "Flavonoids" are a substance or nutrient without an established referenced daily intake value ("RDI").

12. In addition, the package front panel of many Lipton Tea products contains the following statement: "Regular tea drinking, as part of a healthy diet, may help maintain healthy vascular function." Such health claims are in violation of 21 U.S.C. § 352(f)(1), and therefore the products are misbranded.

13. Defendants have made, and continue to make, food label claims that are prohibited by federal and California law. Under federal and California law, Defendants' misbranded food products, including Lipton Tea, cannot legally be manufactured, advertised, distributed, held or sold. Defendants' false and misleading labeling practices stem from their global marketing strategy. Thus, the violations and misrepresentations are similar across product labels and product lines.

### **PARTIES**

14. Plaintiff Amy Maxwell is a resident of San Jose, California who purchased Lipton Tea products in California during the four (4) years prior to the filing of this Complaint (the "Class Period").

15. Defendant Unilever United States, Inc. is a Delaware corporation with its principle place of business at 700 Sylvan Avenue, Englewood Cliffs, New Jersey.

16. Defendant Unilever Supply Chain, Inc. is a Delaware corporation with its principle place of business at 800 Sylvan Avenue, Englewood Cliffs, New Jersey.

17. Defendant PepsiCo, Inc. ("PepsiCo") is a North Carolina corporation with its principle place of business at 700 Anderson Hill Road, Purchase, New York.

1           18. In 1991, Unilever created a joint venture with PepsiCo, the Pepsi Lipton  
2 Partnership (the "Partnership"), for the marketing of ready to drink teas in North America. The  
3 Partnership operates as a subsidiary of PepsiCo, with its principle place of business at 700  
4 Anderson Hill Road, Purchase, New York. PepsiCo and Lipton each control 50% of the shares in  
5 the Partnership. The Partnership manufactures, distributes and sells certain Lipton Tea products.

6           19. Collectively, Defendants are leading producers of retail food products, including  
7 Lipton Tea products. They sell their misbranded food products to consumers through grocery and  
8 other retail stores throughout California.

9                                   **JURISDICTION AND VENUE**

10           20. This Court has original jurisdiction over this action under 28 U.S.C. § 1332(d)  
11 because this is a class action in which: (1) there are over 100 members in the proposed class;  
12 (2) members of the proposed class have a different citizenship from Defendants; and (3) the  
13 claims of the proposed class members exceed \$5,000,000 in the aggregate.

14           21. The Court has jurisdiction over the federal claim alleged herein pursuant to 28  
15 U.S.C. § 1331, because it arises under the laws of the United States.

16           22. The Court has jurisdiction over the California claims alleged herein pursuant to 28  
17 U.S.C. § 1367, because they form part of the same case or controversy under Article III of the  
18 United States Constitution.

19           23. Alternatively, the Court has jurisdiction over all claims alleged herein pursuant to  
20 28 U.S.C. § 1332, because the matter in controversy exceeds the sum or value of \$75,000, and is  
21 between citizens of different states.

22           24. The Court has personal jurisdiction over Defendants because a substantial portion  
23 of the wrongdoing alleged in this Complaint occurred in California, Defendants are authorized to  
24 do business in California, have sufficient minimum contacts with California, and otherwise  
25 intentionally avail themselves of the markets in California through the promotion, marketing and  
26 sale of merchandise, sufficient to render the exercise of jurisdiction by this Court permissible  
27 under traditional notions of fair play and substantial justice.  
28



1           25.     Because a substantial part of the events or omissions giving rise to these claims  
2 occurred in this District and because the Court has personal jurisdiction over Defendants, venue is  
3 proper in this Court pursuant to 28 U.S.C. § 1391(a) and (b).

4                                   **FACTUAL ALLEGATIONS**

5       **A.     Identical California And Federal Laws Regulate Food Labeling**

6           26.     Food manufacturers are required to comply with federal and state laws and  
7 regulations that govern the labeling of food products. First and foremost among these is the  
8 FDCA and its labeling regulations, including those set forth in 21 C.F.R. § 101.

9           27.     Pursuant to the Sherman Law, California has expressly adopted the federal  
10 labeling requirements as its own and indicated that “[a]ll food labeling regulations and any  
11 amendments to those regulations adopted pursuant to the federal act, in effect on January 1, 1993,  
12 or adopted on or after that date shall be the food regulations of this state.” California Health &  
13 Safety Code § 110100.

14           28.     In addition to its blanket adoption of federal labeling requirements, California has  
15 also enacted a number of laws and regulations that adopt and incorporate specific enumerated  
16 federal food laws and regulations. For example, food products are misbranded under California  
17 Health & Safety Code § 110660 if their labeling is false and misleading in one or more  
18 particulars; are misbranded under California Health & Safety Code § 110665 if their labeling fails  
19 to conform to the requirements for nutrient labeling set forth in 21 U.S.C. § 343(q) and  
20 regulations adopted thereto; are misbranded under California Health & Safety Code § 110670 if  
21 their labeling fails to conform with the requirements for nutrient content and health claims set  
22 forth in 21 U.S.C. § 343(r) and regulations adopted thereto; are misbranded under California  
23 Health & Safety Code § 110705 if words, statements and other information required by the  
24 Sherman Law to appear on their labeling are either missing or not sufficiently conspicuous; are  
25 misbranded under California Health & Safety Code § 110735 if they are represented as having  
26 special dietary uses but fail to bear labeling that adequately informs consumers of their value for  
27 that use; and are misbranded under California Health & Safety Code § 110740 if they contain  
28



artificial flavoring, artificial coloring and chemical preservatives but fail to adequately disclose that fact on their labeling.

**B. Defendants' Food Products Are Misbranded**

29. Pursuant to Section 403 of the FDCA, a claim that characterizes the level of a nutrient in a food is a "nutrient content claim" that must be made in accordance with the regulations that authorize the use of such claims. 21 U.S.C. § 343(r)(1)(A). California expressly adopted the requirements of 21 U.S.C. § 343(r) in § 110670 of the Sherman Law.

30. Nutrient content claims are claims about specific nutrients contained in a product. They are typically made on the front of packaging in a font large enough to be read by the average consumer. Because these claims are relied upon by consumers when making purchasing decisions, the regulations govern what claims can be made in order to prevent misleading claims.

31. Section 403(r)(1)(A) of the FDCA governs the use of expressed and implied nutrient content claims on labels of food products that are intended for sale for human consumption. *See* 21 C.F.R. § 101.13.

32. 21 C.F.R. § 101.13 provides the general requirements for nutrient content claims, which California has expressly adopted. *See* California Health & Safety Code § 110100. 21 C.F.R. § 101.13 requires that manufacturers include certain disclosures when a nutrient claim is made and, at the same time, the product contains certain levels of unhealthy ingredients, such as fat and sodium. It also sets forth the manner in which that disclosure must be made, as follows:

(4)(i) The disclosure statement "See nutrition information for \_\_\_ content" shall be in easily legible boldface print or type, in distinct contrast to other printed or graphic matter, and in a size no less than that required by §101.105(i) for the net quantity of contents statement, except where the size of the claim is less than two times the required size of the net quantity of contents statement, in which case the disclosure statement shall be no less than one-half the size of the claim but no smaller than one-sixteenth of an inch, unless the package complies with §101.2(c)(2), in which case the disclosure statement may be in type of not less than one thirty-second of an inch.

(ii) The disclosure statement shall be immediately adjacent to the nutrient content claim and may have no intervening material other than, if applicable, other information in the statement of identity or any other information that is required to be presented with the claim under this section (e.g., see paragraph (j)(2) of this section) or under a regulation in subpart D of this part (e.g., see §§101.54 and 101.62). If the nutrient content claim appears on more than one panel of the label, the disclosure statement shall be adjacent to the claim on each panel except for the

1 panel that bears the nutrition information where it may be omitted.

2 33. An “expressed nutrient content claim” is defined as any direct statement about the  
3 level (or range) of a nutrient in the food (*e.g.*, “low sodium” or “contains 100 calories”). *See* 21  
4 C.F.R. § 101.13(b)(1).

5 34. An “implied nutrient content claim” is defined as any claim that: (i) describes the  
6 food or an ingredient therein in a manner that suggests that a nutrient is absent or present in a  
7 certain amount (*e.g.*, “high in oat bran”); or (ii) suggests that the food, because of its nutrient  
8 content, may be useful in maintaining healthy dietary practices and is made in association with an  
9 explicit claim or statement about a nutrient (*e.g.*, “healthy, contains 3 grams (g) of fat”). 21  
10 C.F.R. § 101.13(b)(2)(i-ii).

11 **1. Defendants Make Unlawful Antioxidant Claims**

12 35. Federal and California regulations regulate antioxidant claims as a particular type  
13 of nutrient content claim. Specifically, 21 C.F.R. § 101.54(g) contains special requirements for  
14 nutrient claims that use the term “antioxidant”:

15 (1) the name of the antioxidant must be disclosed;

16 (2) there must be an established RDI for that antioxidant, and if not, no  
17 “antioxidant” claim can be made about it;

18 (3) the label claim must include the specific name of the nutrient that is an  
19 antioxidant and cannot simply say “antioxidants” (*e.g.*, “high in antioxidant vitamins C and E”),<sup>1</sup>  
20 *see* 21 C.F.R. § 101.54(g)(4);

21 (4) the nutrient that is the subject of the antioxidant claim must also have  
22 recognized antioxidant activity, *i.e.*, there must be scientific evidence that after it is eaten and  
23 absorbed from the gastrointestinal tract, the substance participates in physiological, biochemical  
24

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25  
26 <sup>1</sup> Alternatively, when used as part of a nutrient content claim, the term “antioxidant” or  
27 “antioxidants” (such as “high in antioxidants”) may be linked by a symbol (such as an asterisk)  
28 that refers to the same symbol that appears elsewhere on the same panel of a product label  
followed by the name or names of the nutrients with the recognized antioxidant activity. If this is  
done, the list of nutrients must appear in letters of a type size height no smaller than the larger of  
one half of the type size of the largest nutrient content claim or 1/16 inch.

1 or cellular processes that inactivate free radicals or prevent free radical-initiated chemical  
2 reactions, *see* 21 C.F.R. § 101.54(g)(2);

3 (5) the antioxidant nutrient must meet the requirements for nutrient content claims  
4 in 21 C.F.R. § 101.54(b), (c), or (e) for “High” claims, “Good Source” claims, and “More”  
5 claims, respectively. For example, to use a “High” claim, the food would have to contain 20% or  
6 more of the Daily Reference Value (“DRV”) or RDI per serving. For a “Good Source” claim, the  
7 food would have to contain between 10-19% of the DRV or RDI per serving, *see* 21 C.F.R. §  
8 101.54(g)(3); and

9 (6) the antioxidant nutrient claim must also comply with general nutrient content  
10 claim requirements such as those contained in 21 C.F.R. § 101.13(h) that prescribe the  
11 circumstances in which a nutrient content claim can be made on the label of products high in fat,  
12 saturated fat, cholesterol or sodium.

13 36. The antioxidant labeling for Lipton Tea products violates California law. The  
14 antioxidant claims on the packages of these products violate California law: (1) because the  
15 names of the antioxidants are not disclosed on the product labels; (2) because there are no RDIs  
16 for the antioxidants being touted, including flavonoids; (3) because the claimed antioxidant  
17 nutrients fail to meet the requirements for nutrient content claims in 21 C.F.R. § 101.54(b), (c), or  
18 (e) for “High” claims, “Good Source” claims, and “More” claims, respectively; and (4) because  
19 Defendants lack adequate scientific evidence that the claimed antioxidant nutrients participate in  
20 physiological, biochemical, or cellular processes that inactivate free radicals or prevent free  
21 radical-initiated chemical reactions after they are eaten and absorbed from the gastrointestinal  
22 tract.

23 37. For example, the package label of Lipton Decaffeinated Tea Bags bears the  
24 “Naturally Protective Antioxidants AOX” seal, and the package back panel compares the  
25 “Flavonoid Content” of “Lipton Decaf Tea” to orange juice, broccoli, cranberry juice and coffee.  
26 Additional specific violations are detailed in the FDA Warning Letter, attached hereto as Exhibit  
27 1.  
28

1           38. For these reasons, Defendants' antioxidant claims at issue in this Complaint are  
2 misleading and in violation of 21 C.F.R. § 101.54 and California law, and the products at issue  
3 are misbranded as a matter of law. Misbranded products cannot be legally manufactured,  
4 advertised, distributed, held or sold and are legally worthless.

5           39. In addition to the FDA Warning Letter to Unilever discussed above (Exhibit 1),  
6 the FDA has issued warning letters addressing similar unlawful antioxidant nutrient content  
7 claims. *See, e.g.*, Exhibit 2 (FDA warning letter dated August 30, 2010 to Dr. Pepper Snapple  
8 Group regarding its misbranded Canada Dry Sparkling Green Tea Ginger Ale product because  
9 green tea and green tea flavonoids "are not nutrients with recognized antioxidant activity");  
10 Exhibit 3 (FDA warning letter dated February 22, 2010 to Redco Foods, Inc. regarding its  
11 misbranded Salada Naturally Decaffeinated Green Tea product because "there are no RDIs for  
12 (the antioxidants) grapeskins, rooibos (red tea) and anthocyanins"); Exhibit 4 (FDA warning letter  
13 dated February 22, 2010 to Fleminger Inc. regarding its misbranded TeaForHealth products  
14 because the admonition "[d]rink high antioxidant green tea" . . . "does not include the nutrients  
15 that are the subject of the claim or use a symbol to link the term antioxidant to those nutrients").  
16 Defendants are aware of these FDA warning letters.

17           **2. Defendants Make Unlawful Nutritional Value Claims**

18           40. Defendants have also violated 21 C.F.R. § 101.54(g)(1), which prohibits food  
19 manufacturers from making claims regarding the nutritional value of their products when the  
20 products fail to disclose that no RDI has been established for the touted nutrients.

21           41. For example, certain Lipton Tea products claim to be "packed with flavonoids" but  
22 they fail to disclose that no RDI has been established for flavonoids. Thus, these products violate  
23 21 C.F.R. § 101.54(g)(1).

24           42. The types of misrepresentations made above would be considered by a reasonable  
25 consumer when deciding to purchase Defendants' misbranded food products. The failure to  
26 comply with the labeling requirements of 21 C.F.R. § 101.54 renders Defendants' products  
27 misbranded as a matter of federal and California law.  
28

43. In addition, 21 C.F.R. § 101.65, which has been adopted by California, sets certain minimum nutritional requirements for making an implied nutrient content claim that a product is healthy. For example, for unspecified foods the food must supply at least 10 percent of the RDI of one or more specified nutrients. Defendants have misrepresented the healthiness of their products while failing to meet the regulatory requirements for making such claims.

### **3. Defendants Make Unlawful “Natural” Claims**

44. In its rule-making and warning letters to manufacturers, the FDA has repeatedly stated its policy to restrict the use of the term “natural” in connection with added color, synthetic substances and flavors as provided in 21 C.F.R. § 101.22.

45. The FDA has also repeatedly affirmed its policy regarding the use of the term “natural” as meaning that nothing artificial or synthetic (including all color additives regardless of source) has been included in, or has been added to, a food that would not normally be expected to be in the food.

46. For example, 21 C.F.R. § 70.3(f) makes clear that “where a food substance such as beet juice is deliberately used as a color, as in pink lemonade, it is a color additive.” Similarly, any coloring or preservative can preclude the use of the term “natural” even if the coloring or preservative is derived from natural sources. Further, the FDA distinguishes between natural and artificial flavors in 21 C.F.R. § 101.22.

47. Defendants’ “all natural” labeling practices violate FDA Compliance Guide CPG Sec. 587.100, which states: [t]he use of the words “food color added,” “natural color,” or similar words containing the term “food” or “natural” may be erroneously interpreted to mean the color is a naturally occurring constituent in the food. Since all added colors result in an artificially colored food, we would object to the declaration of any added color as “food” or “natural.”

48. Likewise, California Health & Safety Code § 110740 prohibits the use of artificial flavoring, artificial coloring and chemical preservatives unless those ingredients are adequately disclosed on the labeling.

49. The FDA has sent out numerous warning letters concerning this issue. *See, e.g.*, Exhibit 5 (August 16, 2001 FDA warning letter to Oak Tree Farm Dairy because there was citric

acid in its all natural iced tea); Exhibit 6 (August 29, 2001 FDA warning letter to Hirzel Canning Company because there was citric acid or calcium chloride in its all natural tomato products); Exhibit 7 (August 2, 2001 FDA warning letter to GMP Manufacturing, Inc. stating: “[t]he products, Cytomax Exercise and Recovery Drink (Peachy Keen flavor) and Cytomax Lite (Lemon Iced Tea Flavor) are misbranded because they contain colors but are labeled using the term “no artificial colors.”). Defendants are aware of these FDA warning letters.

50. Defendants have unlawfully labeled a number of their food products as being “all natural” or “made with natural ingredients” when they actually contain artificial ingredients and flavorings, artificial coloring and chemical preservatives. These products include Lipton Pure Leaf Tea products and Lipton Iced Tea products.

51. For example, the label of Lipton 100% Iced Green Tea with Citrus states that it is “100% natural” despite the fact that it contains a number of chemicals, preservatives and artificial flavorings including sugar, citric acid, acerola, fruit extract, REB A (purified stevia extract).

52. Consumers are thus misled into purchasing Defendants’ products with synthetic unnatural ingredients that are not “all natural” as falsely represented on their labeling. Defendants’ products in this respect are misbranded under federal and California law.

#### **4. Defendants Make Unlawful Health Claims**

53. A health claim is a statement expressly or implicitly linking the consumption of a food substance (*e.g.*, ingredient, nutrient, or complete food) to risk of a disease (*e.g.*, cardiovascular disease) or a health-related condition (*e.g.*, hypertension). *See* 21 C.F.R. §101.14(a)(1), (a)(2), and (a)(5). Only health claims made in accordance with FDCA requirements, or authorized by FDA as qualified health claims, may be included in food labeling. Other express or implied statements that constitute health claims, but that do not meet statutory requirements, are prohibited in labeling foods.

54. 21 C.F.R. § 101.14, which has been expressly adopted by California, provides when and how a manufacturer may make a health claim about its product. A “Health Claim” means any claim made on the label or in labeling of a food, including a dietary supplement, that expressly or by implication, including “third party” references, written statements (*e.g.*, a brand

name including a term such as “heart”), symbols (e.g., a heart symbol), or vignettes, characterizes the relationship of any substance to a disease or health-related condition. Implied health claims include those statements, symbols, vignettes, or other forms of communication that suggest, within the context in which they are presented, that a relationship exists between the presence or level of a substance in the food and a disease or health-related condition (see 21 CFR 101.14(a)(1)).

55. Further, health claims are limited to claims about disease risk reduction, and cannot be claims about the diagnosis, cure, mitigation, or treatment of disease. An example of an authorized health claim is: “Three grams of soluble fiber from oatmeal daily in a diet low in saturated fat and cholesterol may reduce the risk of heart disease. This cereal has 2 grams per serving.”

56. A claim that a substance may be used in the diagnosis, cure, mitigation, treatment, or prevention of a disease is a drug claim and may not be made for a food. 21 U.S.C. § 321(g)(1)(D).

57. The use of the term “healthy” is not a health claim but rather an implied nutrient content claim about general nutrition that is defined by FDA regulation. In general, the term may be used in labeling an individual food product that:

Qualifies as both low fat and low saturated fat;

Contains 480 mg or less of sodium per reference amount and per labeled serving, and per 50 g (as prepared for typically rehydrated foods) if the food has a reference amount of 30 g or 2 tbsps or less;

Does not exceed the disclosure level for cholesterol (*e.g.*, for most individual food products, 60 mg or less per reference amount and per labeled serving size); *and*

Except for raw fruits and vegetables, certain frozen or canned fruits and vegetables, and enriched cereal-grain products that conform to a standard of identity, provides at least 10% of the daily value (DV) of vitamin A, vitamin C, calcium, iron, protein, *or* fiber per reference amount. Where eligibility is based on a nutrient that has been added to the food, such fortification must comply with FDA’s fortification policy.



1 21 C.F.R. § 101.65(d)(2). The FDA's definition applies separate criteria to use of healthy on raw,  
2 single ingredient seafood or game meat products. 21 C.F.R. § 101.65(d)(2)(ii). FDA's regulation  
3 on healthy also encompasses other, derivative uses of health (*e.g.*, healthful, healthier) in food  
4 labeling. 21 C.F.R. § 101.65(d).

5 58. Unilever has violated the provisions of § 21 C.F.R. §101.14, 21 C.F.R. §101.65,  
6 21 U.S.C. § 321(g)(1)(D) and 21 U.S.C. § 352(f)(1) on a number of its products and on its  
7 website. For example, until recently on the link to its webpage entitled "Tea and Health,"  
8 subtitled "Heart Health Research" and further subtitled "Cholesterol Research" the following  
9 claim is made: "[F]our recent studies in people at risk for coronary disease have shown a  
10 significant cholesterol lowering effect from tea or tea flavonoids ... One of these studies, on post-  
11 menopausal women, found that total cholesterol was lowered by 8% after drinking 8 cups of  
12 green tea daily for 12 weeks ...." The therapeutic claims on its website establish that the product is  
13 a drug because it is intended for use in the cure, mitigation, treatment, or prevention of disease.  
14 Lipton's products are not generally recognized as safe and effective for the above referenced uses  
15 and, therefore, the product is a "new drug" under section 201(p) of the Act [21 U.S.C. § 321(p)].  
16 New drugs may not be legally marketed in the U.S. without prior approval from FDA as  
17 described in section 505(a) of the Act [21 U.S.C. § 355(a)]. FDA approves a new drug on the  
18 basis of scientific data submitted by a drug sponsor to demonstrate that the drug is safe and  
19 effective.

20 59. As discussed in paragraph 8 and as shown in Exhibit 1, the FDA conducted a  
21 review of one of Defendant's products (Lipton Green Tea 100% Natural Naturally Decaffeinated  
22 Tea) and concluded that Lipton was "in violation of the Federal Food, Drug, and Cosmetic Act ...  
23 and the applicable regulations in Title 21, Code of Federal Regulations, Part 101 (21 CFR 101)."  
24 FDA found the product to be misbranded stating, "Your Lipton Green Tea 100% Natural  
25 Naturally Decaffeinated product is offered for conditions that are not amenable to self-diagnosis  
26 and treatment by individuals who are not medical practitioners; therefore, adequate directions for  
27 use cannot be written so that a layperson can use this drug safely for its intended purposes. Thus,  
28 your Lipton Green Tea 100% Natural Naturally Decaffeinated product is misbranded under

1 section 502(f)(1) of the Act in that the labeling for this drug fails to bear adequate directions for  
2 use [21 U.S.C. § 352(f)(1)].” See Exhibit 1.

3 60. In response to the FDA Warning Letter, Lipton modified its web site and its  
4 packaging by removing some of the most outlandish claims of health and therapeutic benefits that  
5 FDA had found in violation of law. However, there are several objectionable statements on  
6 Lipton’s web site that remain. On the present day web site the following statement appears:  
7 “Flavonoids are dietary compounds found in tea, wine, cocoa, fruit and vegetables. They  
8 contribute significantly to taste and color, and possibly help maintain certain normal, healthy  
9 body functions. A diet rich in flavonoids is generally associated with helping maintain normal,  
10 healthy heart function.” And the package front panel of many Lipton Tea products claims a level  
11 of “flavonoids,” a substance or nutrient without an established referenced daily intake value  
12 (RDI), and contains the following statement, “Regular tea drinking, as part of a healthy diet, may  
13 help maintain healthy vascular function.” Such health claims are in violation of 21 U.S.C. §  
14 352(f)(1) and therefore the products are misbranded.

15 61. Not only do Unilever’s health claims regarding the benefits of “tea flavonoids”  
16 violate FDA rules and regulations, they directly contradict Unilever’s own scientific research,  
17 which has concluded: “[T]he evidence today does not support a direct relationship between tea  
18 consumption and a physiological AOX [antioxidant] benefit.” This conclusion was reported by  
19 Dr. Jane Rycroft, Director of Lipton Tea Institute of Tea, in an article published in January, 2011,  
20 in which Dr. Rycroft states:

21 Only a few scientific publications report an effect of tea on free radical damage in humans  
22 using validated biomarkers in well designed human studies. Unfortunately, the results of  
23 these studies are at variance and the majority of the studies do not report significant  
effects . . .

24 Therefore, despite more than 50 studies convincingly showing that flavonoids possess  
25 potent antioxidant activity *in vitro*, the ability of flavonoids to act as an antioxidant *in vivo*  
[in humans], has not been demonstrated.

26 Based on the current scientific consensus that the evidence today does not support a direct  
relationship between tea consumption and a physiological AOX benefit . . .

27 No evidence has been provided to establish that having antioxidant activity/  
28 content and/or antioxidant properties is a beneficial physiological effect.

1 Rycroft, Jane, "The Antioxidant Hypothesis Needs to be Updated", Vol. 1, *Tea Quarterly Tea*  
2 *Science Overview*, Lipton Tea Institute of Tea Research (Jan. 2011), pp. 2-3. Nonetheless,  
3 Unilever continues to tout the benefits of "tea flavonoids" on its product labels and on its website.

4 **C. Defendants Have Violated California Law**

5 62. Defendants have manufactured, advertised, distributed and sold products that are  
6 misbranded under California law. Misbranded products cannot be legally manufactured,  
7 advertised, distributed or sold and are legally worthless as a matter of law.

8 63. Defendants have violated California Health & Safety Code §§ 109885 and 110390  
9 which make it unlawful to disseminate false or misleading food advertisements that include  
10 statements on products and product packaging or labeling or any other medium used to directly or  
11 indirectly induce the purchase of a food product.

12 64. Defendants have violated California Health & Safety Code § 110395 which makes  
13 it unlawful to manufacture, sell, deliver, hold or offer to sell any misbranded food.

14 65. Defendants have violated California Health & Safety Code § 110398 which makes  
15 it unlawful to deliver or proffer for delivery any food that has been falsely advertised.

16 66. Defendants have violated California Health & Safety Code § 110660 because their  
17 labeling is false and misleading in one or more ways, as follows:

18 a. They are misbranded under California Health & Safety Code § 110665  
19 because their labeling fails to conform to the requirements for nutrient labeling set forth in 21  
20 U.S.C. § 343(q) and the regulations adopted thereto;

21 b. They are misbranded under California Health & Safety Code § 110670  
22 because their labeling fails to conform with the requirements for nutrient content and health  
23 claims set forth in 21 U.S.C. § 343(r) and the regulations adopted thereto; and

24 c. They are misbranded under California Health & Safety Code § 110705  
25 because words, statements and other information required by the Sherman Law to appear on their  
26 labeling either are missing or not sufficiently conspicuous.

67. Defendants have violated California Health & Safety Code § 110760 which makes it unlawful for any person to manufacture, sell, deliver, hold, or offer for sale any food that is misbranded.

68. Defendants have violated California Health & Safety Code § 110765 which makes it unlawful for any person to misbrand any food.

69. Defendants have violated California Health & Safety Code § 110770 which makes it unlawful for any person to receive in commerce any food that is misbranded or to deliver or proffer for deliver any such food.

70. Defendants have violated the standard set by 21 C.F.R. § 101.2, which has been incorporated by reference in the Sherman Law, by failing to include on their product labels the nutritional information required by law.

71. Defendants have violated the standards set by 21 CFR §§ 101.13, and 101.54, which have been adopted by reference in the Sherman Law, by including unauthorized antioxidant claims on their products.

**D. Plaintiff Purchased Defendants' Misbranded Food Products**

72. Plaintiff cares about the nutritional content of food and seeks to maintain a healthy diet.

73. Plaintiff purchased Defendants' misbranded food products at issue in this Complaint, including certain Lipton Tea products, on occasions during the Class Period.

74. Plaintiff purchased the following Lipton Tea products:

Lipton Pureleaf Sweetened bottled iced tea, 16 oz. glass bottle













Lipton Iced Green Tea To Go w/Mandarin & Mango, 14 sticks

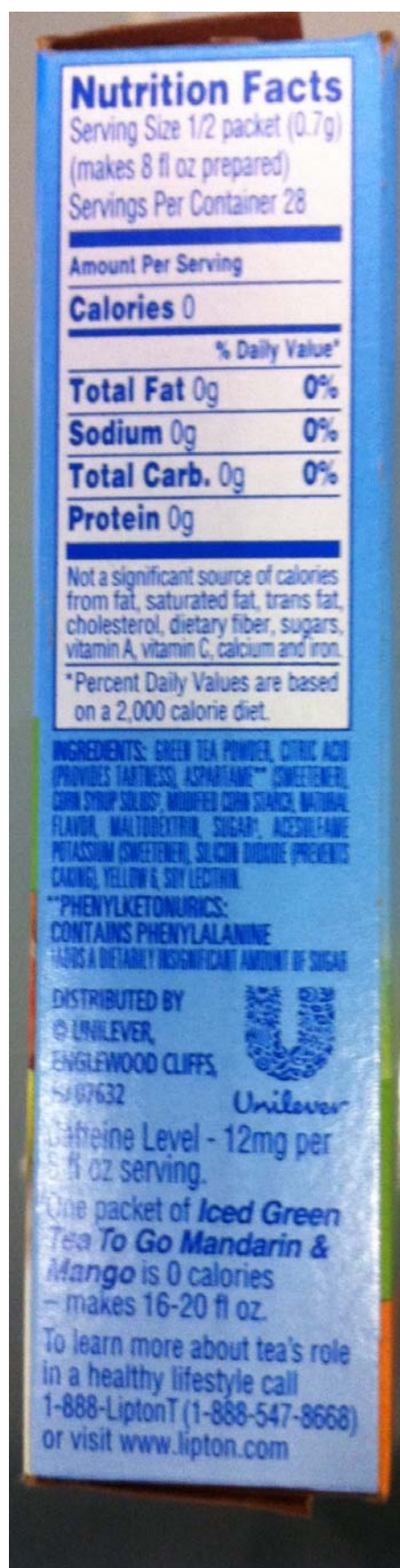




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Lipton Vanilla Caramel Truffle Black Tea, 20 bags









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The advertisement features a central image of a Lipton pyramid tea bag filled with dark tea leaves and small caramel pieces. The bag is set against a warm, orange-gold background. Above the bag, the Lipton logo is prominently displayed in a red shield. To the left of the bag, the words "Long-Leaf Tea" are written in a white, elegant script. To the right, "Real Pieces of Caramel" is written in a similar script. The background is decorated with green tea leaves on the left, and vanilla beans and caramel cubes on the right. Below the main image, the headline "Pyramid Bag – More Room to Infuse." is written in a large, bold, serif font. Underneath the headline, a paragraph of text describes the tea's qualities: "Smooth and enriching, Lipton® Vanilla Caramel Truffle Tea indulges the senses with its satisfying taste and delightful aroma. Our long-cut tea is handpicked from only the top two leaves and a bud and packaged in our unique pyramid-shaped bags that allow the tea to flow freely with real pieces of caramel for a truly authentic tea infusion. Savor the flavor of 100 years' tea expertise in every cup."

**Lipton**

Long-Leaf  
Tea

Real  
Pieces of  
Caramel

**Pyramid Bag – More Room to Infuse.**


Smooth and enriching, Lipton® Vanilla Caramel Truffle Tea indulges the senses with its satisfying taste and delightful aroma. Our long-cut tea is handpicked from only the top two leaves and a bud and packaged in our unique pyramid-shaped bags that allow the tea to flow freely with real pieces of caramel for a truly authentic tea infusion. Savor the flavor of 100 years' tea expertise in every cup.

Lipton Green Tea Decaffeinated, 20 bags






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## Your Small Cup Can Make a Big Difference

Lipton® knows that a great tasting cup of tea starts with the tea leaves. Our **ALL-NATURAL** green tea blend is selected from the freshest high-grown tea leaves and naturally contains **TEA FLAVONOIDS**.

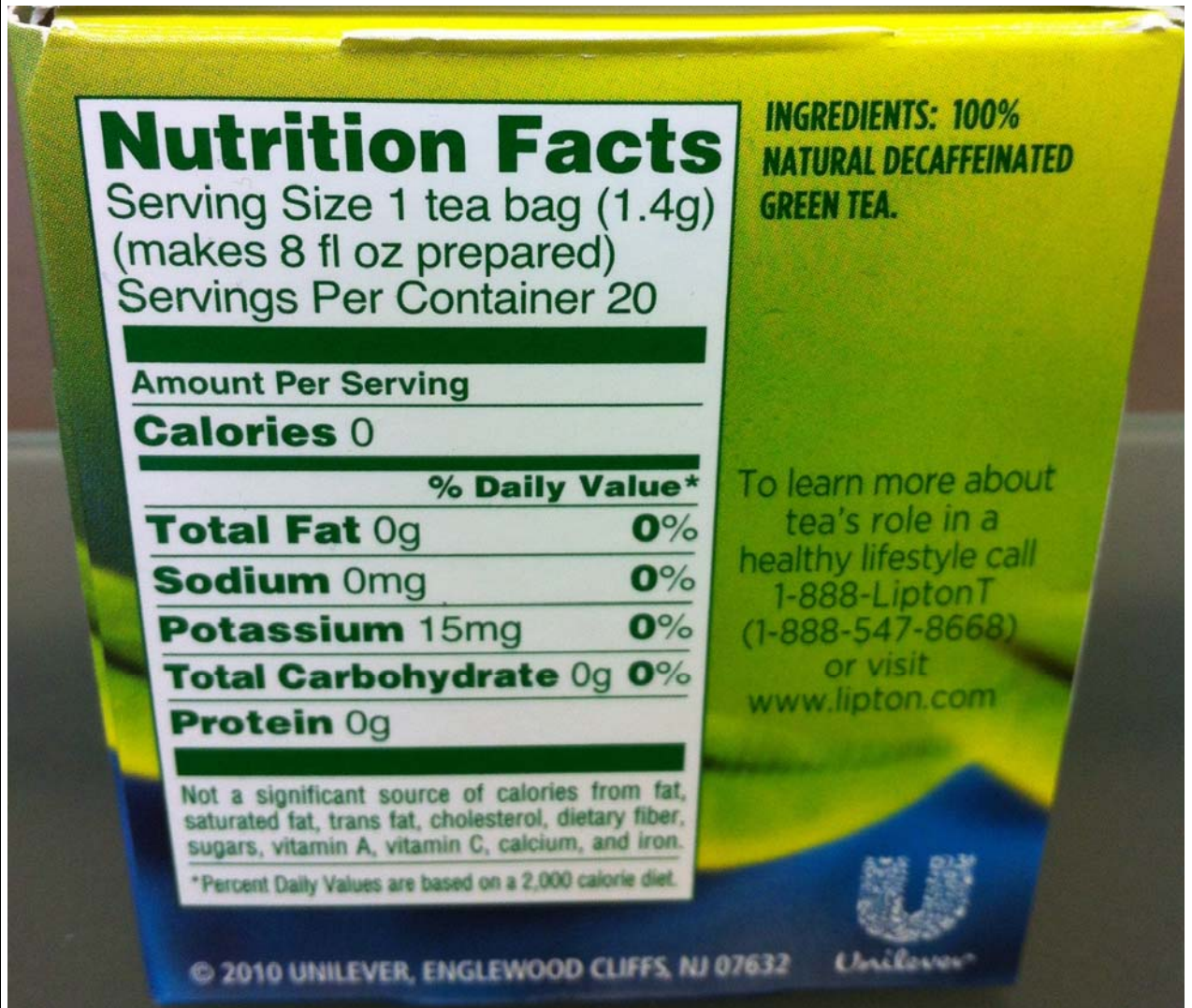
And by drinking a cup of Lipton®, you're not just taking care of yourself. We've sourced our tea from Rainforest Alliance Certified™ tea estates that meet rigorous criteria for protecting the land and providing better conditions for workers, their families and their communities.



*So, go on, drink up... and feel good about the big difference  
your small cup can make.*









Lipton White Tea w/Raspberry, 16.9 oz. plastic bottle



Lipton Cold Brew Iced Tea, 22 bags







Lipton Decaffeinated Tea, 72 bags





**Want to live better everyday?  
Then drink more LIPTON TEA!**

**CONSIDER THESE FACTS:**

Lipton Teas, made from real tea leaves, naturally contain **PROTECTIVE ANTIOXIDANTS** (flavonoid antioxidants.)

This is important because antioxidants help your body protect itself against free radicals-molecules that can damage cells.

**HYDRATION** is key to maintaining a healthy lifestyle. Tea is a great tasting and easy way to get your necessary daily fluid intake!

Lipton Tea is **100% NATURAL**. No additives, preservatives or colorings.

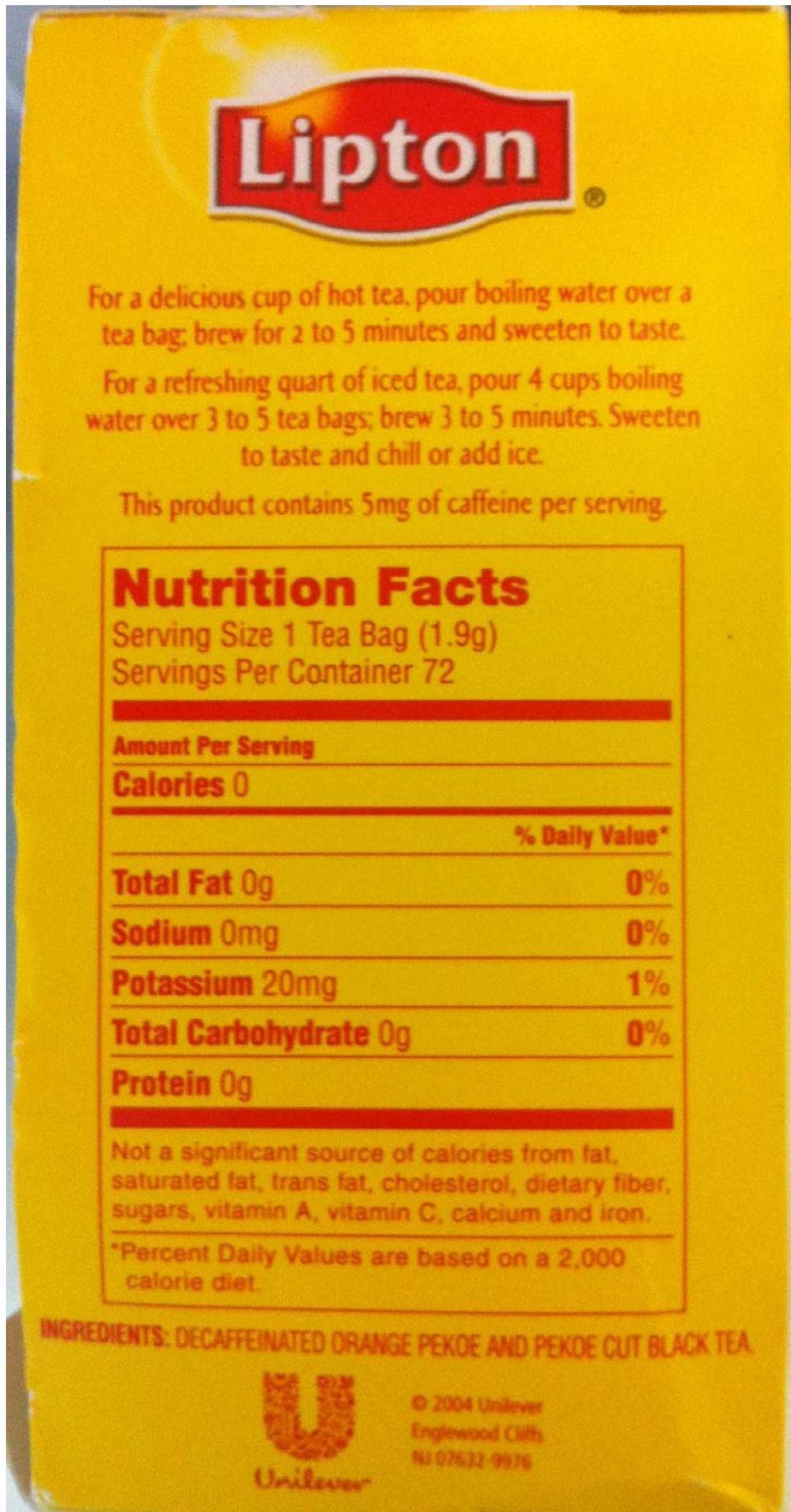
**Flavonoid Content of Selected Beverages & Foods**

Beverage/Food	Flavonoid Content (AOX)
LIPTON DECAF TEA	105
ORANGE JUICE	36
BROCCOLI	10
CRANBERRY JUICE	4
COFFEE	0

Tea is not a substitute for fruits, vegetables and juices.

**Lipton**

*Simply good, naturally protective.*







84. Common Questions Predominate: This action involves common questions of law and fact applicable to each Class member that predominate over questions that affect only individual Class members. Thus, proof of a common set of facts will establish the right of each Class member to recover. Questions of law and fact common to each Class member include, for example:

- a. Whether Defendants engaged in unlawful and misleading business practices by failing to properly package and label their misbranded food products sold to consumers;
- b. Whether the food products at issue were misbranded or unlawfully packaged and labeled as a matter of law;
- c. Whether Defendants made unlawful and misleading antioxidant claims with respect to their food products sold to consumers;
- d. Whether Defendants made unlawful and misleading nutrient content claims with respect to their food products sold to consumers;
- h. Whether Defendants violated California Bus. & Prof. Code § 17200, California Bus. & Prof. Code § 17500, and the Sherman Law;
- i. Whether Plaintiff and the Class are entitled to equitable and/or injunctive relief;
- j. Whether Defendants' unlawful, unfair and/or deceptive practices harmed Plaintiff and the Class; and
- k. Whether Defendants were unjustly enriched by their deceptive practices.

85. Typicality: Plaintiff's claims are typical of the claims of the Class because Plaintiff bought Defendants' misbranded food products during the Class Period. Defendants' unlawful, unfair and/or fraudulent actions concern the same business practices described herein irrespective of where they occurred or were received. Plaintiff and the Class sustained similar injuries arising out of Defendants' conduct in violation of California law. The injuries of each member of the Class were caused directly by Defendants' wrongful conduct. In addition, the factual underpinning of Defendants' misconduct is common to all Class members and represents a common thread of misconduct resulting in injury to all members of the Class. Plaintiff's claims arise from the same practices and course of conduct that give rise to the claims of the Class members and are based on the same legal theories.

1           86.    Adequacy: Plaintiff will fairly and adequately protect the interests of the Class.  
2 Neither Plaintiff nor Plaintiff's counsel have any interests that conflict with or are antagonistic to  
3 the interests of the Class members. Plaintiff has retained highly competent and experienced class  
4 action attorneys to represent their interests and those of the members of the Class. Plaintiff and  
5 Plaintiff's counsel have the necessary financial resources to adequately and vigorously litigate  
6 this class action, and Plaintiff and counsel are aware of their fiduciary responsibilities to the Class  
7 members and will diligently discharge those duties by vigorously seeking the maximum possible  
8 recovery for the Class.

9           87.    Superiority: There is no plain, speedy or adequate remedy other than by  
10 maintenance of this class action. The prosecution of individual remedies by members of the  
11 Class will tend to establish inconsistent standards of conduct for Defendants and result in the  
12 impairment of Class members' rights and the disposition of their interests through actions to  
13 which they were not parties. Class action treatment will permit a large number of similarly  
14 situated persons to prosecute their common claims in a single forum simultaneously, efficiently  
15 and without the unnecessary duplication of effort and expense that numerous individual actions  
16 would engender. Further, as the damages suffered by individual members of the Class may be  
17 relatively small, the expense and burden of individual litigation would make it difficult or  
18 impossible for individual members of the Class to redress the wrongs done to them, while an  
19 important public interest will be served by addressing the matter as a class action. Class  
20 treatment of common questions of law and fact would also be superior to multiple individual  
21 actions or piecemeal litigation in that class treatment will conserve the resources of the Court and  
22 the litigants, and will promote consistency and efficiency of adjudication.

23           88.    The prerequisites to maintaining a class action for injunctive or equitable relief  
24 pursuant to Fed. R. Civ. P. 23(b)(2) are met as Defendants have acted or refused to act on grounds  
25 generally applicable to the Class, thereby making appropriate final injunctive or equitable relief  
26 with respect to the Class as a whole.

27           89.    The prerequisites to maintaining a class action pursuant to Fed. R. Civ. P. 23(b)(3)  
28 are met as questions of law or fact common to class members predominate over any questions

affecting only individual members, and a class action is superior to other available methods for fairly and efficiently adjudicating the controversy.

90. Plaintiff and Plaintiff's counsel are unaware of any difficulties that are likely to be encountered in the management of this action that would preclude its maintenance as a class action.

## **CAUSES OF ACTION**

### **FIRST CAUSE OF ACTION**

#### **Business and Professions Code § 17200, *et seq.* Unlawful Business Acts and Practices**

91. Plaintiff incorporates by reference each allegation set forth above.

92. Defendants' conduct constitutes unlawful business acts and practices.

93. Defendants sold misbranded food products in California during the Class Period.

94. Defendants are corporations and, therefore, each is a "person" within the meaning of the Sherman Law.

95. Defendants' business practices are unlawful under § 17200, *et seq.* by virtue of Defendants' violations of Article 6 (misbranded food) of the Sherman Law.

96. Defendants' business practices are unlawful under § 17200, *et seq.* by virtue of Defendants' violations of § 17500, *et seq.*, which forbids untrue and misleading advertising.

97. Defendants sold Plaintiff and the Class misbranded food products that were not capable of being sold legally and which were legally worthless.

98. As a result of Defendants' illegal business practices, Plaintiff and the Class, pursuant to Business and Professions Code § 17203, are entitled to an order enjoining such future conduct and such other orders and judgments which may be necessary to disgorge Defendants' ill-gotten gains and to restore to any Class Member any money paid for the misbranded food products.

99. Defendants' unlawful business acts present a threat and reasonable continued likelihood of deception to Plaintiff and the Class.